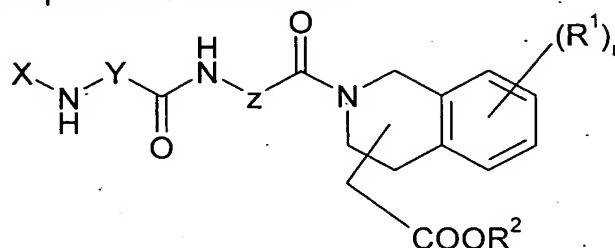


This listing of claims will replace all prior versions, and listings, of claims in the application:

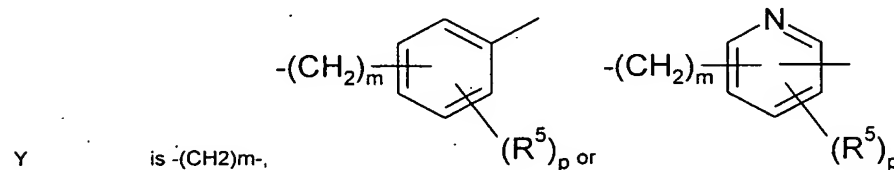
Listing of Claims:

1. (Original) Compounds of the formula I



in which

X is H, $-C(=NR^3)-NHR^4$ or Het,



Y is $-(CH_2)_m-$,

Z is NH or CH_2 ,

R^1

and R^5 are each, independently of one another, H, A, OH, OA, arylalkyl, Hal, $-CO-A$, CN, NO_2 , NHR^3 , COOA, COOH, SO_2A , CF_3 or OCF_3 ,

R^2 is in each case, independently of the others, H or A,

R^3

and R^4 are each, independently of one another, H, A, $-CO-A$, NO_2 or CN,

A is alkyl having 1-6 carbon atoms,

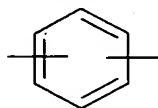
m is 0, 1, 2, 3, 4, 5 or 6,

n and p are, independently of one another, 1, 2 or 3,

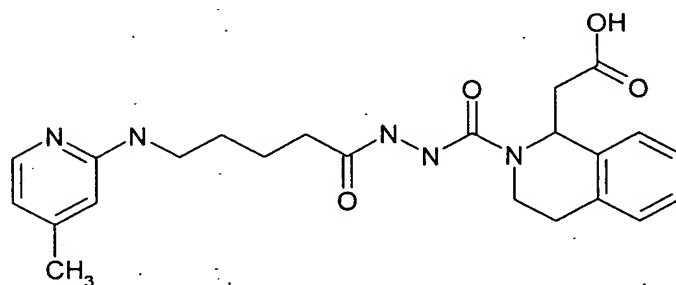
and physiologically acceptable derivatives thereof, in particular salts

and solvates thereof.

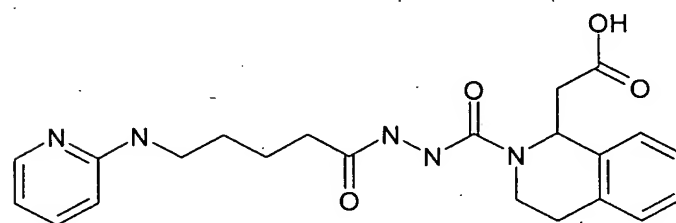
2. (Original) Compounds of the formula I according to Claim 1, in which A is methyl, furthermore ethyl, isopropyl, n-propyl, n-butyl, isobutyl, sec-butyl or tert-butyl.
3. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 and 2~~ Claim 1, in which Het is 4-methylpyridin-2-yl, pyridin-2-yl, pyrimidin-2-yl, imidazol-2-yl, benzimidazol-2-yl and hydrogenated derivatives thereof.
4. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that R^1 and R^5 , independently of one another, are preferably H, A, CN, NO_2 , Hal or -COA.
5. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that R^2 is preferably H or A.
6. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that R^3 and R^4 , independently of one another, are preferably H or -COA.
7. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that X is H, -C(=NH)- NH_2 , -C(=N-methyl)- NH_2 , 4-methylpyridin-2-yl, pyridin-2-yl, pyrimidin-2-yl, imidazol-2-yl, benzimidazol-2-yl and hydrogenated derivatives thereof.
8. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that Y is $-(\text{CH}_2)_m-$ or



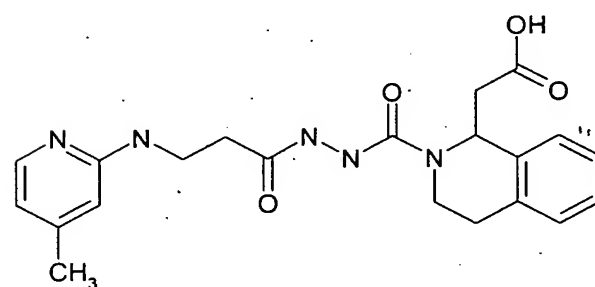
9. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that n and p, independently of one another, are 1 or 2.
10. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 3~~ Claim 1, characterised in that m is 0, 2 or 4.
11. (Original) Compounds of the formulae I1 to I36:



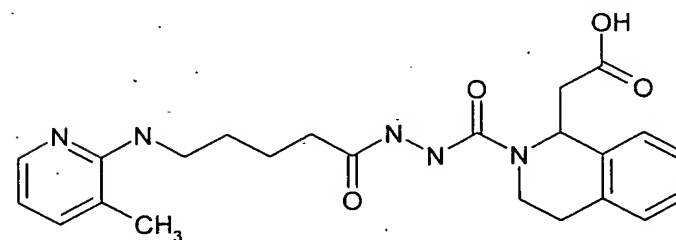
I1



I2

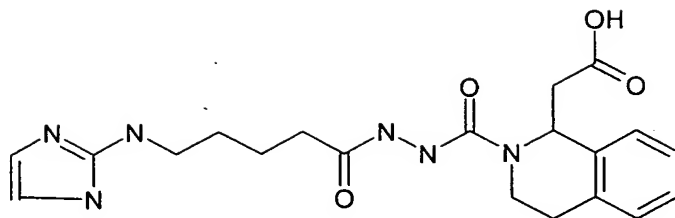


I3

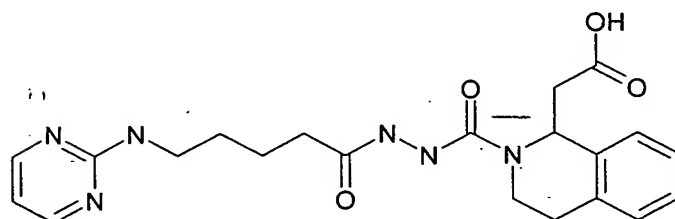


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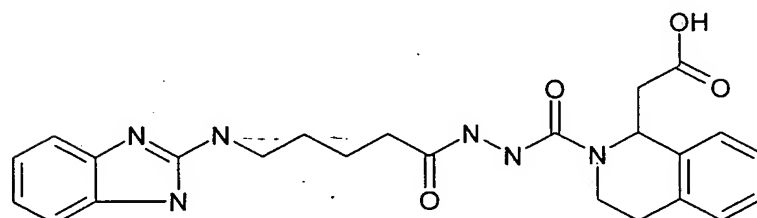
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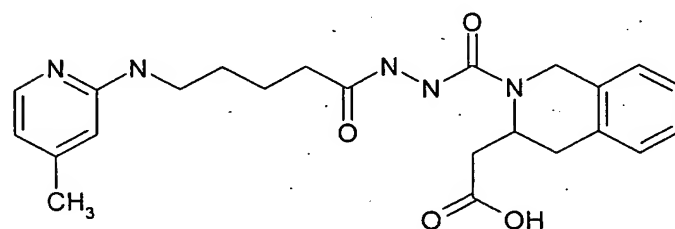
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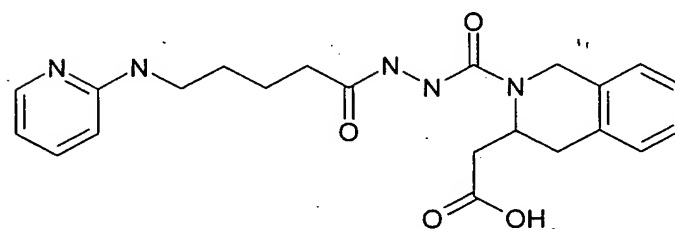
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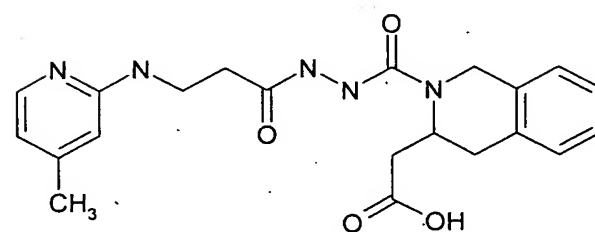
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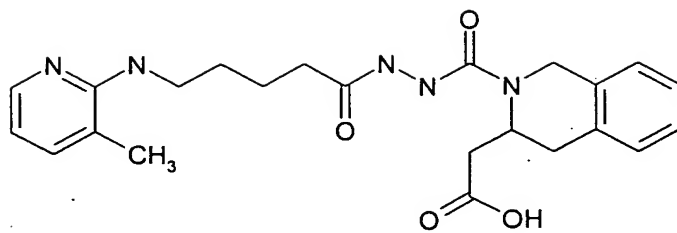
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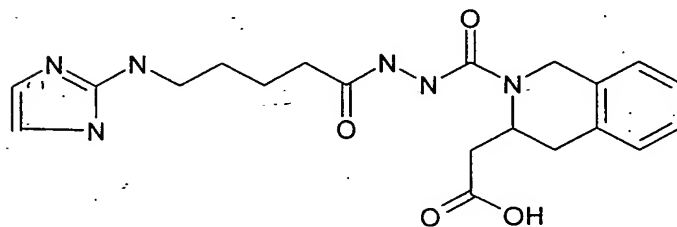
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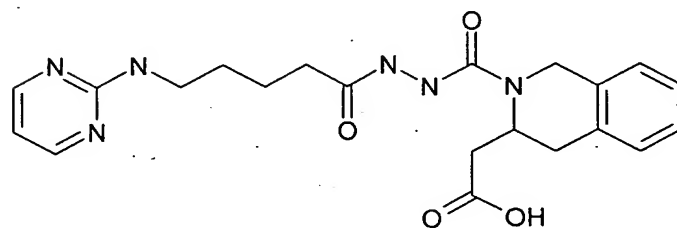
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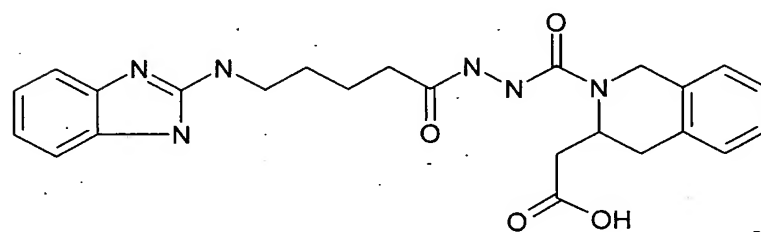
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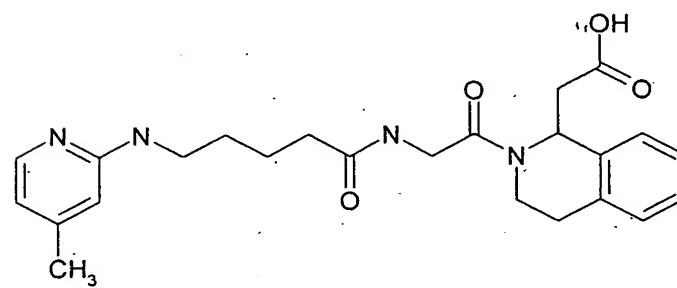
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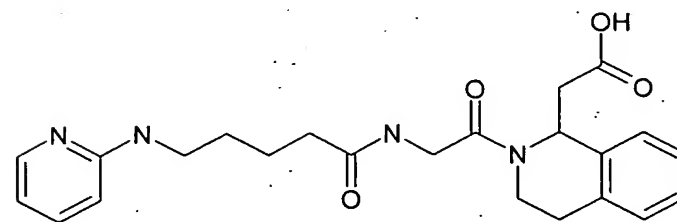
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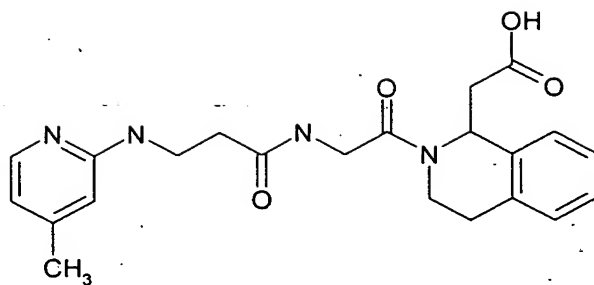
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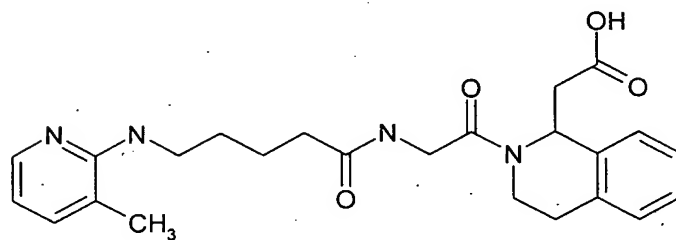
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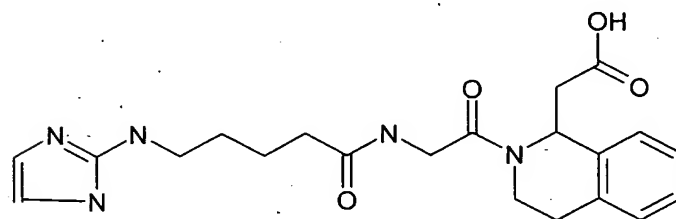
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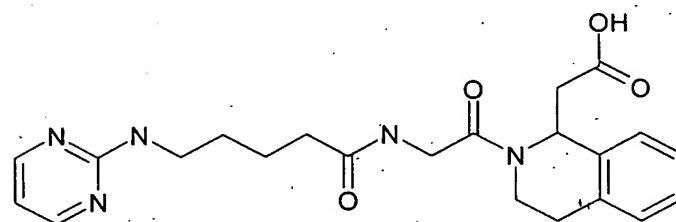
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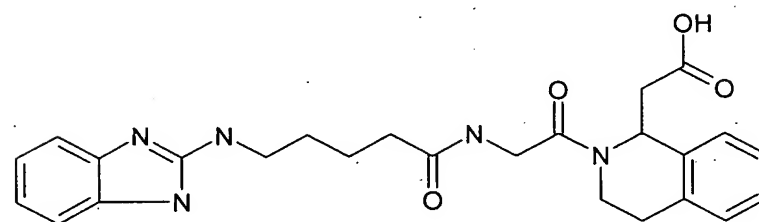
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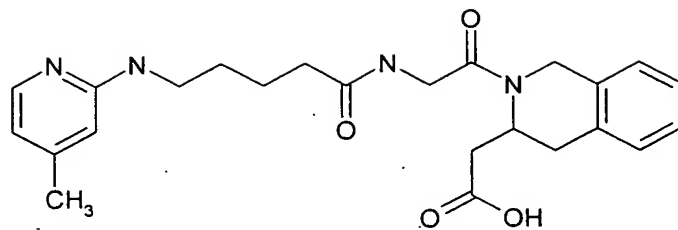
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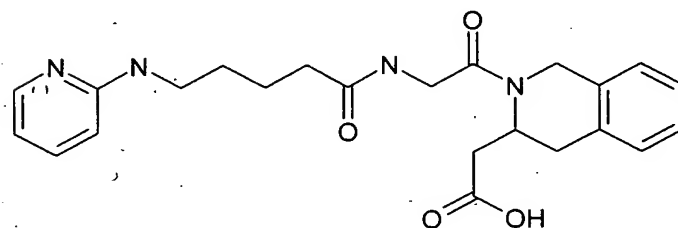
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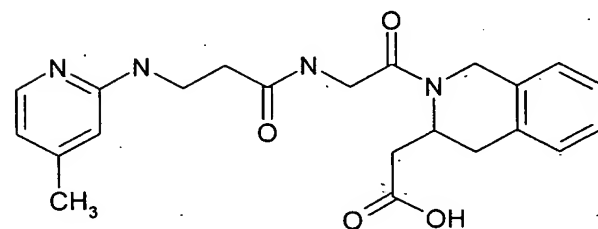
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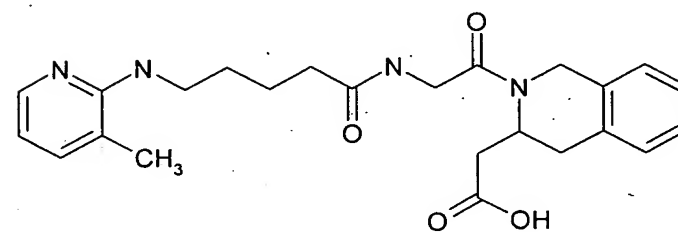
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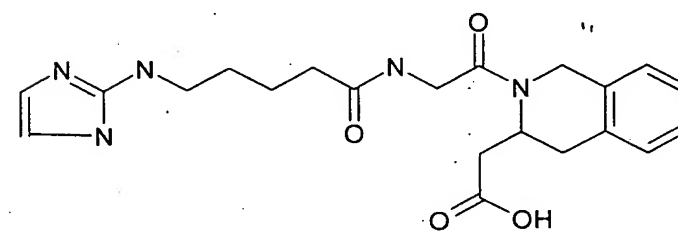
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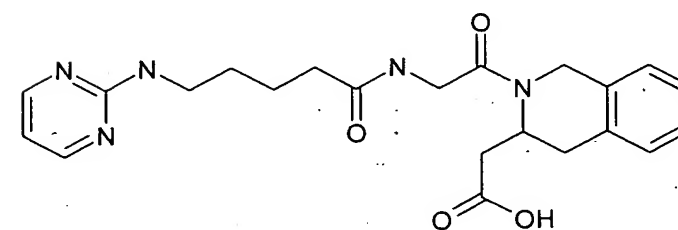
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126

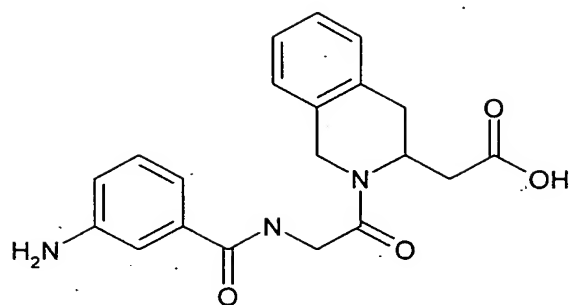


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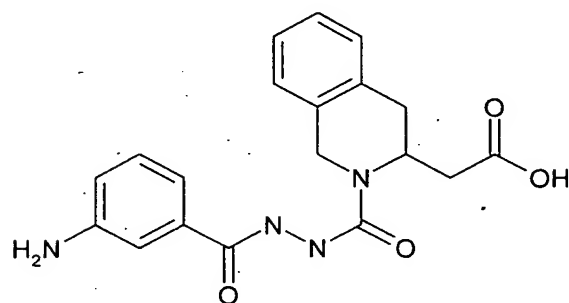


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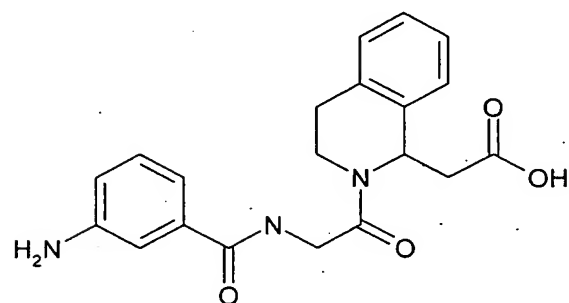
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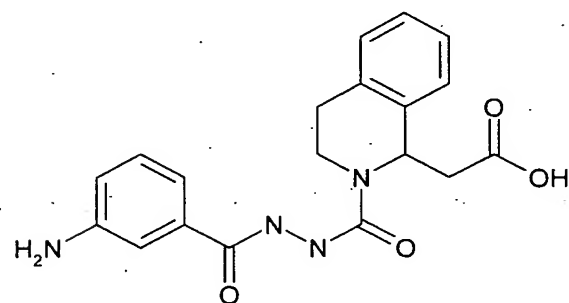
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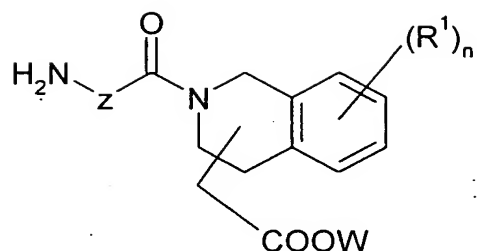
I35



I36



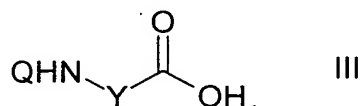
12. (Currently Amended) Process for the preparation of compounds of the formula I according to ~~one or more of Claims 1 to 11~~ Claim 1 and salts thereof, characterised in that
- a) a compound of the formula II



II

in which Z, R¹ and n are as defined above, and W is a conventional protecting group or a solid phase used in peptide chemistry,

is reacted with a compound of the formula III



III

in which Y is as defined above, and Q is a suitable protecting group or Het, in the presence of a condensing agent, such as, for example, HATU,

and the protecting groups and/or the solid phase are subsequently removed,

and, where appropriate, the resultant product is, if Q as protecting group is removed, reacted with a suitable guanyl compound, such as, for example, N,N'-bis-BOC-1-guanylpurazole, and, if desired, the remaining protecting groups and/or the solid phase are removed,

or

b) a compound of the formula I is liberated from one of its functional derivatives by treatment with a solvolysing or hydrogenolysing agent,

and/or in that a basic or acidic compound of the formula I is converted into one of its salts by treatment with an acid or base.

13. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and physiologically acceptable salts or

solvates thereof as therapeutic active ingredients.

14. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and physiologically acceptable salts or solvates thereof as integrin inhibitors.
15. (Currently Amended) Compounds of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and physiologically acceptable salts or solvates thereof for use in combating diseases.
16. (Currently Amended) Pharmaceutical preparation, characterised by a content of at least one compound of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and/or one of its physiologically acceptable salts or solvates.
17. (Currently Amended) Use of compounds of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and/or physiologically acceptable salts or solvates thereof for the preparation of a pharmaceutical preparation.
18. (Currently Amended) Use of compounds of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and/or physiologically acceptable salts or solvates thereof for the preparation of a pharmaceutical preparation for combating thromboses, cardiac infarction, coronary heart diseases, arteriosclerosis, inflammation, tumours, osteoporosis, infections and restenosis after angioplasty.
19. (Currently Amended) Use of compounds of the formula I according to ~~one or more of Claims 1 to 14~~ Claim 1 and/or physiologically acceptable salts or solvates thereof in pathological processes which are maintained or propagated by angiogenesis.